

TITLEPOLYMERIC PHOSPHITE COMPOSITION AND HYDROCYANATION OF
UNSATURATED ORGANIC COMPOUNDS AND THE ISOMERIZATION OF
UNSATURATED NITRILESFIELD OF THE INVENTION

5 This application is a DIV of 09/399,261 filed 09/25/1999, now U.S. Patent 6,284,865
This invention relates to a polymeric phosphite composition and polymeric

phosphite catalyst composition that can be useful for a variety of catalytic
processes, to a process for producing the composition, and to a process for using
the composition in the hydrocyanation of unsaturated organic compounds and the
10 isomerization of unsaturated nitriles.

BACKGROUND OF THE INVENTION

Phosphorus-based ligands are ubiquitous in catalysis, finding use for a
number of commercially important chemical transformations. Phosphorus-based
ligands commonly encountered in catalysis include phosphines and phosphites.

15 Monophosphine and monophosphite ligands are compounds which contain a
single phosphorus atom which serves as a donor to a transition metal.
Bisphosphine, bisphosphite, and bis(phosphorus) ligands in general, contain two
phosphorus donor atoms and typically form cyclic chelate structures with
transition metals.

20 Two industrially important catalytic reactions using phosphorus ligands of
particular importance are olefin hydrocyanation and isomerization of branched
nitriles to linear nitriles. Phosphite ligands are particularly good ligands for both
reactions. The hydrocyanation of ethylenically unsaturated compounds using
transition metal complexes with monodentate phosphite ligands is well
25 documented in the prior art. See for example US 3,496,215; 3,631,191; 3655,723;
3,766,237; and 5,543,536. Bidentate phosphite ligands have also been shown to
be particularly useful ligands in the hydrocyanation of activated ethylenically
unsaturated compounds. See for example, Baker, M.J., and Pringle, P.G., *J.*
Chem. Soc., Chem. Commun., 1292, 1991; Baker et al., *J. Chem. Soc., Chem.*
30 *Commun.*, 803, 1991; WO 93,03839; US 5,512,696; 5,723,641; 5,688,986.

Recovery of the ligand and catalyst is important for a successful process.
Typical separation procedures to remove the product(s) from the catalyst and
ligand involve extraction with an immiscible solvent or distillation. It is usually
difficult to recover the catalyst and ligand quantitatively. For instance, distillation
35 of a volatile product from a non-volatile catalyst results in thermal degradation of
the catalyst. Similarly, extraction results in some loss of catalyst into the product
phase. For extraction, one would like to be able to tune the solubility of the ligand